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on January 13, 2004

*Rimma Mitelman*  
RIMMA MITELMAN  
Reg. No. 34,396  
Attorney for Applicant(s)

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Date of  
Signature

**PATENT**  
#02-0484-UNI  
Case #C4248(C)



**IN THE UNITED STATES PATENT AND TRADEMARK OFFICE**

Applicant: Del Nunzio et al.  
Serial No.: 10/664,398  
Filed: September 17, 2003  
For: Detergent Compositions

Edgewater, New Jersey 07020  
January 13, 2004

**SUBMISSION OF PRIORITY DOCUMENT**

Assistant Commissioner for Patents  
Washington, D.C. 20231

Sir:

Pursuant to rule 55(b) of the Rules of Practice in Patent Cases, Applicant(s) is/are submitting herewith a certified copy of the United Kingdom Application No. 0221740.4 filed September 19, 2002, upon which the claim for priority under 35 U.S.C. § 119 was made in the United States.

It is respectfully requested that the priority document be made part of the file history.

Respectfully submitted,

*Rimma Mitelman*

Rimma Mitelman  
Registration No. 34,396  
Attorney for Applicant(s)

RM/mt  
(201) 840-2671





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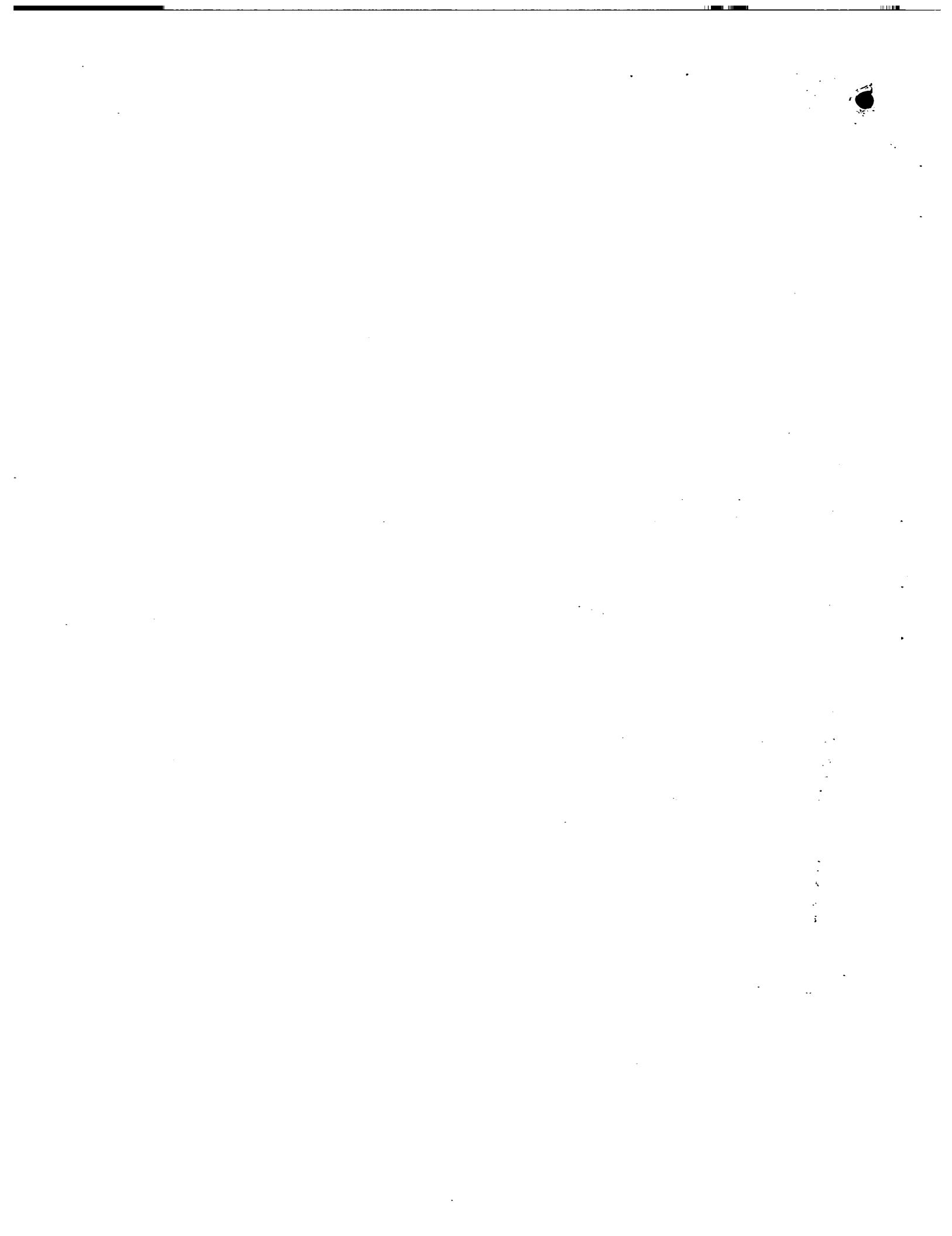
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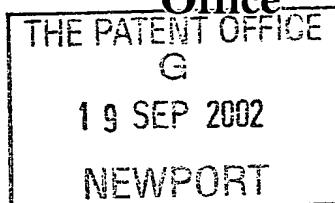
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1. Your reference

C4248(C)/pmk

2. 0221740.4

19 SEP 2002

3. Full name, address and postcode of the or of each applicant (*underline all surnames*)UNILEVER PLC  
UNILEVER HOUSE, BLACKFRIARS  
LONDON, EC4P 4BQPatents ADP number (*if you know it*)

50426956002 ✓

If the applicant is a corporate body, give the country/state of its incorporation

UNITED KINGDOM

4. Title of the invention

DETERGENT COMPOSITIONS

5. Name of your agent (*if you have one*)

ERANSELLA, Mary Evelyn

Elliott, Peter William 30/6/03

"Address for Service" in the United Kingdom to which all correspondence should be sent (*including the postcode*)PATENT DEPARTMENT, UNILEVER PLC  
COLWORTH HOUSE, SHARNBROOK  
BEDFORD, MK44 1LQ

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Patents ADP number (*if you know it*)

6893440501

6573927001

6. If you are declaring priority from one or more earlier patent applications, give the country and the date of filing of the or of each of these earlier applications and (*if you know it*) the or each application numberCountry Priority application number Date of filing  
(*if you know it*) (day / month / year)

7. If this application is divided or otherwise derived from an earlier UK application, give the number and the filing date of the earlier application

Number of earlier application Date of filing  
(day/month/year)8. Is a statement of inventorship and of right to grant of a patent required in support of this request? (*Answer 'Yes' if:*

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- a) any applicant named in part 3 is not an inventor; or
- b) there is an inventor who is not named as an applicant; or
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*See note (d))*

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11. I/We request the grant of a patent on the basis of this application.

Signature(s) 

Date: 18/09/02

Sandra Jane EDWARDS, Authorised Signatory

12. Name and daytime telephone number of person to contact in the United Kingdom **Petra Kimber, Tel 01234 222893**

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DETERGENT COMPOSITIONSTECHNICAL FIELD

5 The present invention relates to effervescent granules for use in particulate detergent compositions.

BACKGROUND AND PRIOR ART

10 Particulate laundry detergent compositions must satisfy a wide range of practical constraints. As well as providing excellent cleaning and care, they must also satisfy numerous physical constraints which go beyond this basic requirement. For example it is known that the effervescence properties of  
15 certain added ingredients can provide improved dispensing qualities to the detergent composition. This can be achieved by adding two ingredients which react together in the presence of water to produce a gas, for example by using sodium carbonate and citric acid. It is also known to place  
20 the two reacting components together in a single granule. Such a single granule is often referred to in the art as an effervescent granule.

US 4 252 664 (Colgate-Palmolive) discloses the use of an  
25 effervescent granule comprising a gas-producing agent (preferably a carbonate) and an acid (preferably a solid acid). The effervescent granule is present in the full formulation at a level of from 5 to 10 wt%.

A series of patent applications have published which disclose effervescent granules which also contain detergent ingredients such as surfactant and builder:

5 WO 98 42811 (Procter & Gamble) discloses a detergent granule of size 1.0 to 4.5 mm which comprises an acid source and an alkali source.

10 WO 98 46714 (Procter & Gamble) discloses a substantially anhydrous effervescent particle of size 0.075 to 20 mm.

15 WO 98 46715 (Procter & Gamble) discloses a substantially anhydrous foaming component comprising an anhydrous surface active component (e.g. surfactant) and an effervescence component (e.g. acid or alkali source).

20 WO 98 46716 (Procter & Gamble) discloses a dry effervescent granule comprising an acid, a carbonate source and a binder, wherein the acid and carbonate source are in close physical proximity.

More recently, it has been discovered that effervescent granules may be improved by controlling the particle sizes of its constituents:

25 WO 00 34422 (Procter & Gamble) discloses an effervescence component which comprises an acid source and a carbonate source wherein at least 75% of the acid source has a particle size of from 0.1 to 150 microns. Preferably the 30 carbonate source also has a defined particle size and the granule is preferably coated in a nonionic surfactant.

WO 01 30949 (Procter & Gamble) discloses a reactive particle comprising two particulate reactants in which the particle number ratio of the first reactant to the second is at least 50:1. Preferably the median particle size of the second to 5 the first reactant is at least 2:1.

The present inventors have surprisingly found that an effervescent granule which comprises a particulate acid source and a particulate carbonate source, wherein the d<sub>50</sub> 10 particle size of the citric acid is greater than, but less than twice as great as, the d<sub>50</sub> particle size of the carbonate source, provides a particularly effective effervescent granule. It is believed that this is due to the inherently greater solubility of the acid source as 15 compared to the carbonate source. The present invention therefore provides an effervescent granule whose constituents are sized such that they dissolve in water at a comparable rate; the inherently more rapid solubility of the acid source being compensated by its increased particle 20 size.

#### STATEMENT OF INVENTION

In a first aspect, the present invention provides an 25 effervescent granule for use in particulate detergent compositions which comprises a first reactant and a second reactant

- (a) a particulate acid source; and
- 30 (b) a particulate carbonate source;

wherein the ratio of the  $d_{50}$  particle size of the acid source to the carbonate source is from less than 2:1 to greater than 1:1.

5 In a second aspect, the present invention provides a laundry detergent composition which comprises from 0.1 to less than 10 wt%, preferably from 0.5 to less than 4 wt%, more preferably from 0.5 to less than 2 wt% of an effervescent granule as defined above.

10

DETAILED DESCRIPTION OF THE INVENTION

The Effervescent Granule

15 The effervescent granule of the present invention comprises a particulate acid source and a particulate carbonate source wherein the  $d_{50}$  particle size of the citric acid is greater than, but less than twice as great as, the  $d_{50}$  particle size of the carbonate source. Without wishing to be bound by theory, it is believed that this may be because the acid source is inherently more rapidly dissolving than the carbonate source. This means that when particles of the same size are added to water under the same conditions, the particles of acid dissolve more rapidly. Hence the increased particle size of the acid source slows down its solubility such that the solubility of the carbonate source becomes more comparable and an enhanced effervescent action results.

- 5 -

Preferably, the ratio of the  $d_{50}$  particle size of the acid source to the carbonate source is from 1.9:1 to 1.1:1, more preferably from 1.8:1 to 1.2:1.

5 The granule preferably comprises from 20 to 80 wt%, preferably from 30 to 60 wt%, more preferably from 40 to 50 wt% of the carbonate source. The granule preferably comprises from 10 wt% to 60 wt%, preferably from 20 wt% to 50 wt%, more preferably from 30 wt% to 40 wt% of the acid  
10 source.

It is preferred that the majority of the granule is comprised of the carbonate and the acid source and that more preferably the carbonate source and the acid source make up  
15 from 50 to 100 wt%, preferably from 60 to 99 wt%, more preferably from 70 to 95 wt%, most preferably from 80 to 90 wt% of the granule.

The granules preferably have a particle size of from 0.2 to  
20 10 mm, preferably from 0.5 to 5 mm, more preferably 0.8 to 4 mm, most preferably from 1 to 2 mm. This helps to provide a highly effective localised fizzing action.

25 The granule may also comprise a drying agent such as a silica gel to ensure that the granule remains dry during storage.

30 The granule may also comprise a binder to help hold it together and it is preferred that such a binder is a polyethylene glycol or a polypropylene glycol.

The granules preferably also comprises a colourant in order to contrast their appearance from that of the detergent compositions to which they are added.

5    The Acid Source

The acid source reacts with the carbonate source in the presence of water to produce carbon dioxide gas.

10   Preferably the acid source is a particulate material and has a narrow particle size distribution such that at least 50 wt%, preferably at least 60 wt%, more preferably at least 70 wt%, most preferably at least 80 wt% of particles have a particle size of from 150 to 350 microns.

15   Preferably the acid source is an organic acid. Preferably the acids are mono-, bi- or tri-protonic acids. Such preferred acids include mono- or polycarboxylic acids preferably citric acid, adipic acid, glutaric acid, 20 citramalic acid, tartaric acid, maleic acid, malic acid, succinic acid, malonic acid. Such acids are preferably used in their acidic forms, and it may be preferred that their anhydrous forms are used, or mixtures thereof. Citric acid is the most highly preferred.

25   The Carbonate Source

The carbonate source reacts with the acid source in the presence of water to produce carbon dioxide gas.

The carbonate source is preferably selected from carbonate, bicarbonate, sesquicarbonate and mixtures thereof.

5 The carbonate source may advantageously be a combination of sodium bicarbonate and sodium carbonate in a weight ratio of greater than 1:1, preferably from greater than 1:1 to 100:1, preferably from greater than 3:2 to 75:1, more preferably from 2:1 to 50:1, most preferably from 3:1 to 25:1.

10 The Optional Solid Surfactant Particles

In order to enhance the fizzing action and to provide a positive cue for the end user, the effervescent granules of the present invention preferably comprise solid surfactant particles. It is preferred that in its isolated state the surfactant is sufficiently solid such that it is pourable.

As desired, the granule may comprises from 1 to 15 wt%, preferably from 2 to 10 wt%, more preferably from 3 to 8 wt% 20 of solid surfactant particles.

Preferably the surfactant has a melting point of at least 30°C, preferably at least 40°C, more preferably at least 50°C.

25 Surfactants suitable for the present invention meet the criteria of being solid and particulate. Highly preferred surfactants are the alkyl sulphate type surfactants, preferably a C<sub>8</sub>-C<sub>15</sub> primary alkyl sulphate, more preferably a

C<sub>10</sub>-C<sub>14</sub> primary alkyl sulphate. Also preferred are the C<sub>8</sub>-C<sub>15</sub> olefin sulphonates.

Preferably the granules are substantially free of alkyl  
5 benzene sulphonate surfactants, because of their  
hygroscopicity. In this context, 'substantially free' means  
less than 5 wt% of the effervescent granule, preferably less  
than 3 wt%, more preferably less than 1 wt%, most preferably  
completely absent.

10

Preferably the surfactant particles have a d<sub>50</sub> particle size  
of greater than 100 microns, more preferably from 150 to  
800 microns.

15 In a particularly preferred embodiment, the ratio of the  
particle size of the effervescent granule to the d<sub>50</sub>  
particle size of the surfactant is greater than 5:1,  
preferably from 6:1 to 1000:1.

20 The Method of Manufacture

The effervescent granules according to the present invention  
may be made by any suitable process, such as for example by  
granulation, roll-compaction, extrusion or a combination  
25 thereof.

The Detergent Composition

The effervescent granules of the present invention are  
30 intended to be added to a conventional particulate laundry

detergent composition. When added to a detergent composition they are present at a level of from 0.1 to less than 10 wt%, preferably from 0.5 to less than 4 wt%, more preferably from 0.5 to less than 2 wt%. The benefit of the  
5 effervescent granules is surprisingly not diminished at reduced levels of inclusion.

Detergent compositions of the invention contain detergent-active compounds and detergency builders, and may optionally  
10 contain bleaching components and other active ingredients to enhance performance and properties.

Detergent-active compounds (surfactants) may be chosen from soap and non-soap anionic, cationic, nonionic, amphoteric  
15 and zwitterionic detergent-active compounds, and mixtures thereof. Many suitable detergent-active compounds are available and are fully described in the literature, for example, in "Surface-Active Agents and Detergents", Volumes I and II, by Schwartz, Perry and Berch. The preferred  
20 detergent-active compounds that can be used are soaps and synthetic non-soap anionic and nonionic compounds. The total amount of surfactant present is suitably within the range of from 5 to 40 wt%.

25 Anionic surfactants are well-known to those skilled in the art. Examples include alkylbenzene sulphonates, particularly linear alkylbenzene sulphonates having an alkyl chain length of C<sub>8</sub>-C<sub>15</sub>; primary and secondary alkylsulphates, particularly C<sub>8</sub>-C<sub>15</sub> primary alkyl sulphates;  
30 alkyl ether sulphates; olefin sulphonates; alkyl xylene

sulphonates; dialkyl sulphosuccinates; and fatty acid ester sulphonates. Sodium salts are generally preferred.

Nonionic surfactants that may be used include the primary 5 and secondary alcohol ethoxylates, especially the C<sub>8</sub>-C<sub>20</sub> aliphatic alcohols ethoxylated with an average of from 1 to 20 moles of ethylene oxide per mole of alcohol, and more especially the C<sub>10</sub>-C<sub>15</sub> primary and secondary aliphatic alcohols ethoxylated with an average of from 1 to 10 moles 10 of ethylene oxide per mole of alcohol. Non-ethoxylated nonionic surfactants include alkylpolyglycosides, glycerol monoethers, and polyhydroxyamides (glucamide).

Cationic surfactants that may be used include quaternary 15 ammonium salts of the general formula R<sub>1</sub>R<sub>2</sub>R<sub>3</sub>R<sub>4</sub>N<sup>+</sup> X<sup>-</sup> wherein the R groups are long or short hydrocarbyl chains, typically alkyl, hydroxyalkyl or ethoxylated alkyl groups, and X is a solubilising cation (for example, compounds in which R<sub>1</sub> is a C<sub>8</sub>-C<sub>22</sub> alkyl group, preferably a C<sub>8</sub>-C<sub>10</sub> or C<sub>12</sub>-C<sub>14</sub> alkyl 20 group, R<sub>2</sub> is a methyl group, and R<sub>3</sub> and R<sub>4</sub>, which may be the same or different, are methyl or hydroxyethyl groups); and cationic esters (for example, choline esters).

Detergent compositions suitable for use in most automatic 25 fabric washing machines generally contain anionic non-soap surfactant, or nonionic surfactant, or combinations of the two in any ratio, optionally together with cationic, amphoteric or zwitterionic surfactants, optionally together with soap.

The detergent compositions of the invention also contain one or more detergency builders. The total amount of detergency builder in the compositions will suitably range from 5 to 80 wt%, preferably from 10 to 60 wt%.

5

Suitable inorganic non-phosphorus containing builders include crystalline and amorphous aluminosilicates, for example zeolites as disclosed in GB 1 473 201 (Henkel); amorphous aluminosilicates as disclosed in GB 1 473 202 10 (Henkel); and mixed crystalline/amorphous aluminosilicates as disclosed in GB 1 470 250 (Henkel); and layered silicates as disclosed in EP 164 514B. Inorganic phosphate builders, for example, sodium orthophosphate, pyrophosphate and tripolyphosphate, may also be present.

15

Aluminosilicates, include the zeolite used in most commercial particulate detergent compositions, namely zeolite A. Advantageously, however, maximum aluminium zeolite P (zeolite MAP) described and claimed in EP 384 070B 20 (Unilever) may be used. Zeolite MAP is an alkali metal aluminosilicate of the P type having a silicon to aluminium ratio not exceeding 1.33, preferably not exceeding 1.15, and more preferably not exceeding 1.07.

25 Suitably the builder system comprises a zeolite (for example zeolite A) and optionally an alkali metal citrate and/or a crystalline layered silicate (for example SKS-6 ex Hoechst).

30 Examples of phosphorus-containing inorganic detergency builders include the water-soluble salts, especially the

alkali metal salts of pyrophosphates, orthophosphates, polyphosphates and phosphonates.

The phosphorus containing inorganic builder is preferably  
5 pyrophosphate or polyphosphate. Specific examples of  
inorganic phosphate builders include sodium and potassium  
tripolyphosphates, orthophosphates and hexametaphosphates.

Organic builders that may be present include polycarboxylate  
10 polymers such as polyacrylates and acrylic/maleic  
copolymers; monomeric polycarboxylates such as citrates,  
gluconates, oxydisuccinates, glycerol mono-, di- and  
trisuccinates, carboxymethyloxysuccinates,  
carboxymethyloxymalonates, dipicolinates,  
15 hydroxyethyliminodiacetates, alkyl- and alkenylmalonates and  
succinates; and sulphonated fatty acid salts. These  
materials are preferably present in alkali metal salt,  
especially sodium salt, form. This list is not intended to  
be exhaustive.

Especially preferred organic builders are citrates, suitably  
used in amounts of from 5 to 30 wt%, preferably from 10 to  
25 wt%; and acrylic polymers, more especially  
acrylic/maleic copolymers, suitably used in amounts of from  
25 0.5 to 15 wt%, preferably from 1 to 10 wt%. Builders, both  
inorganic and organic, are preferably present in alkali  
metal salt, especially sodium salt, form.

Detergent compositions according to the invention may also  
30 suitably contain a bleach system. Preferably this will  
include a peroxy bleach compound, for example, an inorganic

persalt or an organic peroxyacid, capable of yielding hydrogen peroxide in aqueous solution. Preferred inorganic persalts are sodium perborate monohydrate and tetrahydrate, and sodium percarbonate. The sodium percarbonate may have a protective coating against destabilisation by moisture. The peroxy bleach compound is suitably present in an amount of from 5 to 35 wt%, preferably from 10 to 25 wt%.

The peroxy bleach compound may be used in conjunction with a bleach activator (bleach precursor) to improve bleaching action at low wash temperatures. The bleach precursor is suitably present in an amount of from 1 to 8 wt%, preferably from 2 to 5 wt%. Preferred bleach precursors are peroxycarboxylic acid precursors, more especially peracetic acid precursors and peroxybenzoic acid precursors; and peroxycarbonic acid precursors. An especially preferred bleach precursor suitable for use in the present invention is N,N,N',N'-tetracetyl ethylenediamine (TAED).

A bleach stabiliser (heavy metal sequestrant) may also be present. Suitable bleach stabilisers include ethylenediamine tetraacetate (EDTA), diethylenetriamine pentaacetate (DTPA), ethylenediamine disuccinate (EDDS), and the polyphosphonates such as the Dequests (Trade Mark), ethylenediamine tetramethylene phosphonate (EDTMP) and diethylenetriamine pentamethylene phosphate (DETPMP).

Bleach ingredients are postdosed.

The compositions of the invention may contain alkali metal, preferably sodium, carbonate, in order to increase

detergency and ease processing. Sodium carbonate may suitably be present in amounts ranging from 1 to 60 wt%, preferably from 2 to 40 wt%. Sodium carbonate may be included in either or both base granule, and/or may be 5 postdosed.

As previously indicated, sodium silicate may also be present. The amount of sodium silicate may suitably range from 0.1 to 5 wt%. Sodium silicate may be postdosed, for 10 example, as granular sodium disilicate, or as sodium carbonate/sodium silicate cogranules, for example, Nabion (Trade Mark) 15 ex Rhodia Chimie.

Powder flow may be improved by the incorporation of a small 15 amount of a powder structurant. Examples of powder structurants, some of which may play other roles in the formulation as previously indicated, include, for example, fatty acids (or fatty acid soaps), sugars, acrylate or acrylate/maleate polymers, sodium silicate, and dicarboxylic 20 acids (for example, Sokalan (Trade Mark) DCS ex BASF). One preferred powder structurant is fatty acid soap, suitably present in an amount of from 1 to 5 wt%.

Other materials that may be present in detergent 25 compositions of the invention include antiredeposition agents such as cellulosic polymers; soil release agents; anti-dye-transfer agents; fluorescers; inorganic salts such as sodium sulphate; enzymes (proteases, lipases, amylases, cellulases); dyes; coloured speckles; perfumes; 30 and fabric conditioning compounds. This list is not intended to be exhaustive.

EXAMPLES

An effervescent granule was made with the following composition:

5

Ingredient	Wt%
Citric acid	38
Sodium bicarbonate	42
Sodium carbonate	10
Sodium C <sub>12-14</sub> alkyl sulphate	5
Polypropylene glycol	4.5
Silica Gel	0.5
Total	100

The sodium carbonate had a d<sub>50</sub> particle size of about 150 microns. The sodium bicarbonate had a d<sub>50</sub> particle size of 10 about 120 microns. The citric acid had a d<sub>50</sub> particle size of about 250 microns. The sodium C<sub>12-C14</sub> alkyl sulphate was in particulate form (with a melting point in excess of 60°C) and had a d<sub>50</sub> particle size of about 200 microns.

15 The sodium carbonate, sodium bicarbonate, silica gel, sodium lauryl sulphate and citric acid were mixed together to provide a dry-mix. Polypropylene glycol was then added to produce a wetted mixture. The resultant mixture was then granulated in a sieve-type granulator with 2 mm holes. The 20 extrudates were then sliced into cylinders with a length of approximately 2 mm.

1 parts of the effervescent granule was mixed with 99 parts of a phosphate detergent powder with a formulation given below:

Ingredient	Wt%
<b>Base Powder</b>	
Sodium C <sub>12</sub> alkyl benzene sulphonate	18.0
Sodium tripolyphosphate	12.0
Sodium silicate	7.5
Sodium sulphate	26.9
SCMC	0.50
Optical brightener	0.18
Photobleach	0.03
Silicone	0.02
Salts + moisture	0.5
<b>Post Dosed</b>	
Sodium carbonate	15.0
Sodium sulphate	10.0
Protease	0.24
Amylase	0.10
Sodium perborate monohydrate	1.50
TAED	0.50
Perfume	0.24
Moisture	6.8
<b>Total</b>	<b>100.00</b>

The resultant detergent powder gave an attractive effervescent action when added to wash water despite the very low level of effervescent granule.

CLAIMS

1. An effervescent granule for use in particulate detergent compositions which comprises

5

(a) a particulate acid source; and  
(b) a particulate carbonate source;

wherein the ratio of the  $d_{50}$  particle size of the acid source to the carbonate source is from less than 2:1 to greater than 1:1.

10 2. An effervescent granule as claimed in claim 1, wherein the ratio of the  $d_{50}$  particle size of the acid source to the carbonate source is from 1.9:1 to 1.1:1, preferably from 1.8:1 to 1.2:1.

15 3. An effervescent granule as claimed in any preceding claim, wherein the effervescent granules comprise from 10 wt% to 60 wt%, preferably from 20 wt% to 50 wt%, more preferably from 30 wt% to 40 wt% of the acid source.

20 4. An effervescent granule as claimed in any preceding claim, wherein the effervescent granules comprise from 20 to 80 wt%, preferably from 30 to 60 wt%, more preferably from 40 to 50 wt% of the carbonate source.

25 5. An effervescent granule as claimed in any preceding claim, wherein the effervescent granule comprises at

from 50 to 100 wt%, preferably from 60 to 99 wt%, more preferably from 70 to 95 wt%, most preferably from 80 to 90 wt%, of the total of the acid source and carbonate source.

5

6. An effervescent granule as claimed in any preceding claim, which is substantially free of alkyl benzene sulphonate surfactant.

10 7. An effervescent granule as claimed in any preceding claim, wherein the acid source is citric acid.

15 8. A laundry detergent composition which comprises from 0.1 to less than 10 wt%, preferably from 0.5 to less than 4 wt%, more preferably from 0.5 to less than 2 wt% of an effervescent granule as claimed in any one of claims 1 to 7.

20